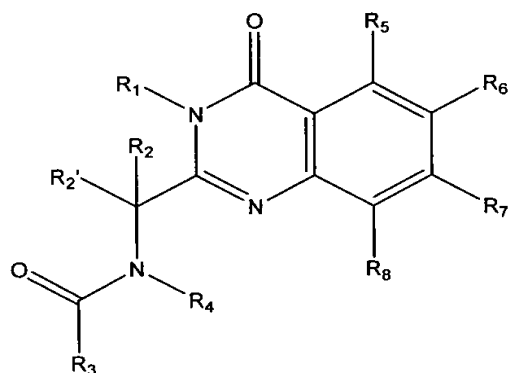


In the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Previously Amended) A method of treating cellular proliferative diseases comprising administering a compound chosen from the group consisting of:



wherein:

R₁ is chosen from hydrogen, alkyl, aryl, alkylaryl, heteroaryl, alkylheteroaryl, substituted alkyl, substituted aryl, substituted alkylaryl, substituted heteroaryl, and substituted alkylheteroaryl;

R₂ and R₂' are independently chosen from hydrogen, alkyl, oxaalkyl, aryl, alkylaryl, heteroaryl, alkylheteroaryl, substituted alkyl, substituted aryl, substituted alkylaryl, substituted heteroaryl, and substituted alkylheteroaryl; or R₂ and R₂' taken together form a 3- to 7-membered ring;

R₃ is chosen from hydrogen, alkyl, aryl, alkylaryl, heteroaryl, alkylheteroaryl, substituted alkyl, substituted aryl, substituted alkylaryl, substituted heteroaryl, substituted alkylheteroaryl, oxaalkyl, oxaalkylaryl, substituted oxaalkylaryl, R₁₅O- and R₁₅-NH-;

R₄ is chosen from alkyl, aryl, alkylaryl, alkylheteroaryl, substituted alkyl, and substituted aryl;

R₅, R₆, R₇ and R₈ are independently chosen from hydrogen, alkyl, alkoxy, halogen, fluoroalkyl, nitro, dialkylamino, alkylsulfonyl, alkylsulfonamido, sulfonamidoalkyl, sulfonamidoaryl, alkylthio, carboxyalkyl, carboxamido, aminocarbonyl, aryl and heteroaryl; and

R₁₅ is chosen from alkyl, aryl, alkylaryl, heteroaryl, alkylheteroaryl, substituted alkyl, substituted aryl, substituted alkylaryl, substituted heteroaryl, and substituted alkylheteroaryl; or a pharmaceutically acceptable salt of any of the foregoing compounds.

2. (Cancelled)

3. (Cancelled)

4. (Previously amended) A method according to claim 1 wherein

R₁ is chosen from hydrogen, alkyl, aryl, substituted alkyl, substituted aryl, heteroaryl, substituted heteroaryl, alkylaryl, substituted alkylaryl and substituted alkylheteroaryl;

R₂ is chosen from hydrogen, alkyl and substituted alkyl;

C1 R₂' is hydrogen;

R₃ is chosen from alkyl, substituted alkyl, alkylaryl, heteroaryl, aryl, substituted aryl, substituted heteroaryl, substituted oxaalkylaryl R₁₅O- and R₁₅-NH-;

R₄ is chosen from alkyl, aryl, alkylaryl, alkylheteroaryl, substituted alkyl, and substituted aryl;

R₅ is hydrogen;

R₆, R₇ and R₈ are independently chosen from hydrogen, halogen, methyl and trifluoromethyl; and

R₁₅ is chosen from alkyl, aryl and substituted aryl.

5. (Cancelled)

6. (Cancelled)

7. (Previously Amended) A method according to claim 4 wherein R₁ is chosen from hydrogen, lower alkyl, substituted lower alkyl, benzyl, substituted benzyl, phenyl, naphthyl and substituted phenyl.

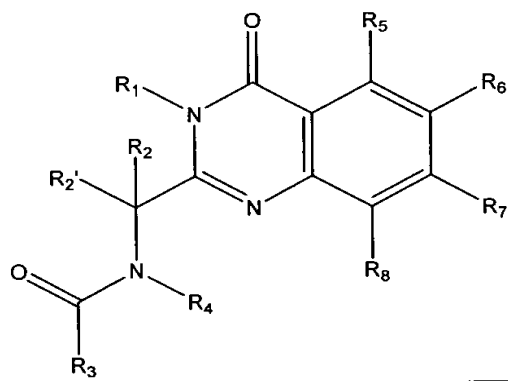
8. - (Original) A method according to claim 7 wherein R₁ is chosen from hydrogen, ethyl, propyl, methoxyethyl, naphthyl, phenyl, bromophenyl, chlorophenyl, methoxyphenyl, ethoxyphenyl, tolyl, dimethylphenyl, chlorofluorophenyl, methylchlorophenyl, ethylphenyl, phenethyl, benzyl, chlorobenzyl, methylbenzyl, methoxybenzyl, tetrahydrofuranylmethyl and (ethoxycarbonyl)ethyl.

9. (Previously Amended) A method according to claim 4 wherein R₂ is chosen from hydrogen, lower alkyl and substituted lower alkyl, and R₂' is hydrogen.

10. (Original) A method according to claim 9 wherein R₂ is chosen from hydrogen, methyl, ethyl, propyl, methylthioethyl, aminobutyl, (CBZ)aminobutyl, cyclohexylmethyl, benzyloxymethyl, methylsulfinylethyl, methylsulfinylmethyl, hydroxymethyl, benzyl and indolylmethyl.

11. (Previously Amended) A method according to claim 4 wherein R₃ is chosen from C₁-C₁₃ alkyl; substituted lower alkyl; phenyl; naphthyl; phenyl substituted with one or more halo, lower alkyl, loweralkoxy, nitro, carboxy, methylenedioxy or trifluoromethyl; biphenyl; benzyl; phenoxyethyl; halophenoxyethyl; phenylvinyl; heteroaryl; heteroaryl substituted with lower alkyl; and benzyloxymethyl.

12. (Currently Amended) **A method of treating cellular proliferative diseases comprising administering a compound:**



R₁ is chosen from hydrogen, alkyl, aryl, substituted alkyl, substituted aryl, heteroaryl, substituted heteroaryl, alkylaryl, substituted alkylaryl and substituted alkylheteroaryl;

R₂ is chosen from hydrogen, alkyl and substituted alkyl;

R₂' is hydrogen;

~~A method according to claim 11 wherein~~

R₃ is chosen from ethyl, propyl, chloropropyl, butoxy, heptyl, butyl, octyl, tridecanyl, (ethoxycarbonyl)ethyl, dimethylaminoethyl, dimethylaminomethyl, phenyl, naphthyl,

halophenyl, dihalophenyl, cyanophenyl, halo(trifluoromethyl)phenyl, chlorophenoxymethyl, methoxyphenyl, carboxyphenyl, ethylphenyl, tolyl, biphenyl, methylenedioxyphenyl, methylsulfonylphenyl, methoxychlorophenyl, chloronaphthyl, methylhalophenyl, trifluoromethylphenyl, butylphenyl, pentylphenyl, methylnitrophenyl, phenoxymethyl, dimethoxyphenyl, phenylvinyl, nitrochlorophenyl, nitrophenyl, dinitrophenyl, bis(trifluoromethyl)phenyl, benzyloxymethyl, benzyl, furanyl, benzofuranyl, pyridinyl, indolyl, methylpyridinyl, quinoliny, picoliny, pyrazolyl, and imidazolyl;

R₄ is chosen from alkyl, aryl, alkylaryl, alkylheteroaryl, substituted alkyl, and substituted aryl;

R₅ is hydrogen; and

R₆, R₇ and R₈ are independently chosen from hydrogen, halogen, methyl and trifluoromethyl;

or a pharmaceutically acceptable salt of any of the foregoing compounds.

13. (Previously Amended) A method according to claim 4 wherein R₃ is R₁₅-NH- and R₁₅ is chosen from lower alkyl; cyclohexyl; phenyl; and phenyl substituted with halo, lower alkyl, loweralkoxy, or lower alkylthio.

14. (Currently Amended) A method according to claim ~~13~~ **4** wherein R₁₅ is chosen from isopropyl, butyl, cyclohexyl, phenyl, bromophenyl, dichlorophenyl, methoxyphenyl, ethylphenyl, tolyl, trifluoromethylphenyl and methylthiophenyl.

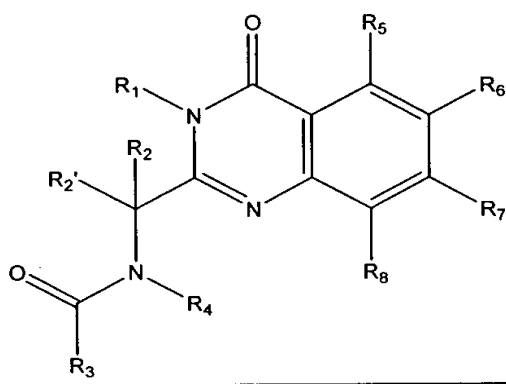
15. (Previously Amended) A method according to claim 4 wherein R₄ is chosen from lower alkyl, substituted lower alkyl, cyclohexyl; phenyl substituted with hydroxy, lower alkoxy or lower alkyl; benzyl; heteroarylmethyl; heteroarylethyl; and heteroarylpropyl.

16. (Currently Amended) A method according to claim ~~4~~ **15** wherein R₄ is chosen from methyl, ethyl, propyl, butyl, cyclohexyl, carboxyethyl, carboxymethyl, methoxyethyl, hydroxyethyl, hydroxypropyl, dimethylaminoethyl, dimethylaminopropyl, diethylaminoethyl, diethylaminopropyl, aminopropyl, methylaminopropyl, 2,2-dimethyl-3-(dimethylamino)propyl, 1-cyclohexyl-4-(diethylamino)butyl, aminoethyl, aminobutyl, aminopentyl, aminoethyl, aminoethoxyethyl, isopropylaminopropyl, diisopropylaminoethyl, 1-methyl-4-(diethylamino)butyl, (t-Boc)aminopropyl, hydroxyphenyl, benzyl, methoxyphenyl,

methoxymethoxyphenyl, dimethylphenyl, tolyl, ethylphenyl, (oxopyrrolidinyl)propyl, (methoxycarbonyl)ethyl, benzylpiperidinyl, pyridinylethyl, pyridinylmethyl, morpholinylethyl, morpholinylpropyl, piperidinyl, azetidinylmethyl, azetidinypropyl, pyrrolidinylethyl, pyrrolidinylpropyl, piperidinylmethyl, piperidinylethyl, imidazolylpropyl, imidazolylethyl, (ethylpyrrolidinyl)methyl, (methylpyrrolidinyl)ethyl, (methylpiperidinyl)propyl, (methylpiperazinyl)propyl, furanylmethyl and indolylethyl.

17. (Currently Amended) ~~A method according to claim 4~~

A method of treating cellular proliferative diseases comprising administering a compound:



wherein

R₁ is chosen from lower alkyl, benzyl, substituted benzyl and substituted phenyl;

R₂ is chosen from hydrogen, alkyl, substituted lower alkyl and benzyl;

R₂' is hydrogen;

R₃ is chosen from substituted phenyl and naphthyl;

R₄ is substituted alkyl;

R₅ is hydrogen or halo

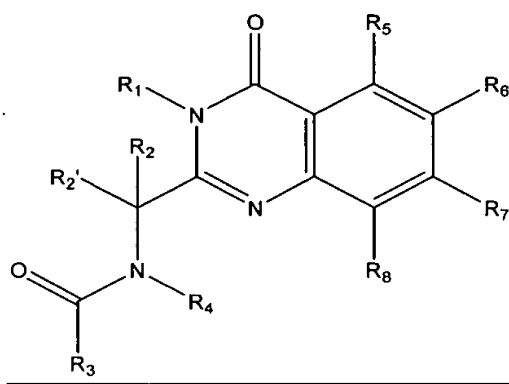
R₆ is hydrogen, methyl or halo;

R₇ is hydrogen, halo, methyl or trifluoromethyl; and

R₈ is hydrogen or halo;

or a pharmaceutically acceptable salt of any of the foregoing compounds.

18. (Currently Amended) **A method of treating cellular proliferative diseases comprising administering a compound having the structure of:**



~~A method according to claim 1~~ wherein

R₁ is benzyl or halobenzyl;

R₂ is chosen from ethyl and propyl;

R₂' is hydrogen;

R₃ is substituted phenyl;

R₄ is (CH₂)_m OH or (CH₂)_p R₁₆ wherein m is 2 or 3 and p is 1-3;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is halo;

R₈ is hydrogen;

R₁₆ is chosen from amino, propylamino, and azetidiny;

or a pharmaceutically acceptable salt of any of the foregoing compounds.

19. (Original) A method according to claim 18 wherein the stereogenic center to which R₂ and R₂' are attached is of the R configuration.

20-29. (Cancelled)

30. (Currently Amended) A method according to claim 1, ~~or 2~~ wherein said disease or disorder is chosen from the group consisting of cancer, hyperplasia, restenosis, cardiac hypertrophy, immune disorders and inflammation.

31-59. (Cancelled)

60. (Withdrawn)

61. (Withdrawn)

62. (Withdrawn)

63. (Previously Added) The method of claim 1, wherein:

R₁ is benzyl or halobenzyl;

R₂ is ethyl or propyl;

R₂' is hydrogen;

R₃ is substituted phenyl;

R₄ is $-(CH_2)_mOH$ or $-(CH_2)_pR_{16}$ wherein m is two or three and p is one to three;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is halo;

R₈ is hydrogen; and

R₁₆ is chosen from amino, propylamino, and azetidiny;

or a pharmaceutically acceptable salt thereof.

64. (Previously Added) The method of claim 1, wherein wherein R₃ is phenyl substituted with one or more halo, lower alkyl, loweralkoxy, nitro, carboxy, methylenedioxy, or trifluoromethyl.

65. (Previously Added) The method of claim 1, wherein

R₁ is benzyl;

R₂ is isopropyl;

R₂' is hydrogen;

R₃ is p-tolyl;

R₄ is 3-aminopropyl;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is chloro; and

R₈ is hydrogen.

66. (Previously Added) The method of claim 1, wherein said salt is a mesylate.

67. (Previously Added) The method of claim 1, wherein

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(isopropylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is p-chlorobenzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(dimethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is m-methoxybenzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

C1 R₁ is benzyl; R₂ is isopropyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is azetidin-3-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-aminoethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-aminoethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(methylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(methylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(methylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is azetidin-2-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is methylsulfinylmethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is piperidin-3-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is fluoro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-aminoethyl; R₅, R₆, R₇ and R₈ are hydrogen;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is piperidin-2-yl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 4-aminobutyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

C I R₁ is m-chlorobenzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(piperidin-1-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(imidazol-3-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is pyrrolidin-3-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(diethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-chlorophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 4-aminobutyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is pyrrolidin-2-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(azetidin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(pyrrolidin-1-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(pyrrolidin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(dimethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is propyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(pyrrolidin-1-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(pyrrolidin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is piperidin-4-ylmethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is methylsulfinylethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(piperidin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is benzyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is (N-ethylpyrrolidin-2-yl)methyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-piperidinyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 4-piperidinyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is p-chlorobenzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2,2-dimethyl-3-(dimethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 5-aminopentyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-(dimethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is fluoro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(2-methylpiperidin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is fluoro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(N-methylpyrrolidin-2-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-trifluoromethylphenyl; R₄ is 3-(dimethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(diethylamino)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 3-(N-methylpiperazin-1-yl)propyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is 4-(CBZ)aminobutyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

C1 R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is aminoethoxyethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is 2-naphthyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro

R₁ is benzyl; R₂ is cyclohexylmethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(piperidin-1-yl)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-hydroxypropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-fluorophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 6-aminoethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₇, and R₈ are hydrogen; and R₆ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is fluoro;

R₁ is benzyl; R₂ is methyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-aminoethyl; R₅, R₆, R₇ and R₈ are hydrogen;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₇ are hydrogen; and R₈ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₆, R₇, and R₈ are hydrogen; and R₅ is chloro;

R₁ is benzyl; R₂ is aminobutyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 3-aminopropyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-tolyl; R₄ is 2-(dimethylamino)ethyl; R₅ and R₈ are hydrogen; and R₆ and R₇ are fluoro;

R₁ is m-tolyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro;

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-(dimethylamino)ethyl; R₅ and R₈ are hydrogen; and R₆ and R₇ are fluoro; or

R₁ is benzyl; R₂ is ethyl; R₂' is hydrogen; R₃ is p-bromophenyl; R₄ is 2-carboxyethyl; R₅, R₆, and R₈ are hydrogen; and R₇ is chloro,

or a pharmaceutically acceptable salt of any of the foregoing compounds.

68. (New) A method according to claim 1 wherein the stereogenic center to which R₂ and R₂' are attached is of the R configuration.

69. (New) A method according to claim 12 wherein the stereogenic center to which R₂ and R₂' are attached is of the R configuration.

70. (New) A method according to claim 17 wherein the stereogenic center to which R₂ and R₂' are attached is of the R configuration.

71. (New) A method according to claim 12 wherein said disease or disorder is chosen from the group consisting of cancer, hyperplasia, restenosis, cardiac hypertrophy, immune disorders and inflammation.

72. (New) A method according to claim 17 wherein said disease or disorder is chosen from the group consisting of cancer, hyperplasia, restenosis, cardiac hypertrophy, immune disorders and inflammation.

73. (New) A method according to claim 18 wherein said disease or disorder is chosen from the group consisting of cancer, hyperplasia, restenosis, cardiac hypertrophy, immune disorders and inflammation.

74. (New) The method of claim 17, wherein:

R₁ is benzyl or halobenzyl;

R₂ is ethyl or propyl;

R₂' is hydrogen;

R₃ is substituted phenyl;

R₄ is $-(CH_2)_mOH$ or $-(CH_2)_pR_{16}$ wherein m is two or three and p is one to three;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is halo;

R₈ is hydrogen; and

R₁₆ is chosen from amino, propylamino, and azetidiny;

or a pharmaceutically acceptable salt thereof.

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75. (New) The method of claim 17, wherein wherein R₃ is phenyl substituted with one or more halo, lower alkyl, loweralkoxy, nitro, carboxy, methylenedioxy, or trifluoromethyl.

76. (New) The method of claim 18, wherein wherein R₃ is phenyl substituted with one or more halo, lower alkyl, loweralkoxy, nitro, carboxy, methylenedioxy, or trifluoromethyl.

77. (New) The method of claim 12, wherein

R₁ is benzyl;

R₂ is isopropyl;

R₂' is hydrogen;

R₃ is p-tolyl;

R₄ is 3-aminopropyl;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is chloro; and

R₈ is hydrogen.

78. (New) The method of claim 17, wherein

R₁ is benzyl;

R₂ is isopropyl;

R₂' is hydrogen;

R₃ is p-tolyl;

R₄ is 3-aminopropyl;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is chloro; and

R₈ is hydrogen.

79. (New) The method of claim 18, wherein

R₁ is benzyl;

R₂ is isopropyl;

R₂' is hydrogen;

R₃ is p-tolyl;

R₄ is 3-aminopropyl;

R₅ is hydrogen;

R₆ is hydrogen;

R₇ is chloro; and

R₈ is hydrogen.

80. (New) The method of claim 12, wherein said salt is a mesylate.

81. (New) The method of claim 17, wherein said salt is a mesylate.

82. (New) The method of claim 18, wherein said salt is a mesylate.
